## What is claimed is:

- 1. A biocompatible, supplemented tissue sealant composition comprising:
  (i) at least one antibody, and (ii) fibrinogen in an amount which is capable of forming a fibrin matrix in the presence of thrombin, Factor XIII and Ca<sup>++</sup>.
- 2. The supplemented tissue sealant composition of claim 1, wherein said antibody is released long term.
- 3. The supplemented tissue sealant composition of claim 2, wherein said antibody interacts with said fibrin matrix.
- 4. The supplemented tissue sealant composition of claim 2, wherein antibody is of sufficiently low solubility to permit localized, sustained-release of antibody.
- 5. The supplemented tissue scalant composition of claim 2, wherein said antibody is in solid form.
- 6. The supplemented tissue sealant composition of claim 5, wherein said antibody is introduced into said matrix in solution in a carrier, said carrier having a higher rate of dissolution than said composition contained therein, so that the composition is deposited within the matrix as a solid precipitate.
- 7. The supplemented tissue sealant composition of claim 2, wherein the mass of antibody exceeds an amount which is soluble in the volume of the matrix, thereby permitting localized, sustained-release of antibody.
- 8. The supplemented tissue sealant composition of claim 7, wherein the antibody is introduced into said matrix as an emulsion.

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- 9. The supplemented tissue sealant composition of elaims 1 or 2, wherein said antibody is selected from the group consisting of polyclonal antibodies, monoclonal antibodies, chimeric antibodies, recombinant antibodies, transgenic antibodies, functional derivatives and fragments thereof, including modified forms thereof.
- The supplemented tissue scalant composition of claims 1 or 2, further comprising an effective amount of at least one additional supplement selected from the group consisting of analgesics, anticogulants, anti-inflammatory compounds, antimicrobial compositions, antiproliferatives, cytotoxins, chemotherapeutic drugs, cytokines, growth factors, hormones, interferons, lipids, oligonucleotides, osteoinducers, polymers, polysaccharides, protease inhibitors, proteoglycans, polypeptides, steroids, vasoconstrictors, vasodilators, vitamins, minerals and stabilizers.
- 11. The supplemented tissue sealant composition of claim 10, wherein said polysaccharide supplement is selected from the group consisting of chitin, chitosan and derivatives thereof.
- 12. An antibody delivery system comprising a biocompatible, supplemented tissue sealant composition, said composition comprising: (i) at least one antibody, and (ii) fibrinogen in an amount which is capable of forming a fibrin matrix in the presence of thrombin, Factor XIII and Ca<sup>++</sup>.
- 13. The delivery system of claim 12, wherein the supplemented tissue sealant composition is delivered to tissue of a patient, thereby permitting the localized release of said antibody to the tissue of the patient.
- Q 14. The delivery system of claims 12 or 13, wherein said localized release of the antibody is sustained release.

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- 15. The delivery system of claims 14, wherein the antibody is of sufficiently low solubility to permit localized, sustained-release of antibody.
- 16. The delivery system of claims 14, wherein the mass of antibody exceeds an amount which is soluble in the volume of the matrix, thereby permitting localized, sustained-release of antibody.
- 17. The delivery system of claim 16, wherein said antibody is introduced into said matrix as an emulsion.
- 18. The delivery system of claims 14, wherein said antibody interacts with said fibrin matrix, thereby permitting localized, sustained-release of said antibody.
- 19. The delivery system of claim 14, wherein said antibody is in solid form.
- 20. The delivery system of claim 19, wherein said antibody is introduced into said matrix in solution in a carrier, said carrier having a higher rate of dissolution than said composition contained therein, so that the composition is deposited within the matrix as a solid precipitate.

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